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(71) Applicant: CORNING INCORPORATED
Corning, N.Y. 14831 (US)

(72) Inventors:

• Berkey, George Edward
SP FR 02-12, Corning, NY 14831 (US)

• Liou Wan-I, Lisa
SP FR 02-12, Corning, NY 14831 (US)
• Modavis, Robert Adam
SP FR 02-12, Corning, NY 14831 (US)
• Nolan, Daniel Aloysius
SP FR 02-12, Corning, NY 14831 (US)
• Weldman, David Lee
SP FR 02-12, Corning, NY 14831 (US)

(74) Representative:

Marchant, James Ian et al
Elkington and Fife,
Prospect House,
8 Pembroke Road
Sevenoaks, Kent TN13 1XR (GB)

(54) Athermalized codoped optical waveguide device

(57) Athermalized optical waveguide devices and methods of making the athermalized devices are described. Boron is incorporated into the composition of the optical waveguides in order to athermalize the waveguides by reducing spectral shifts caused by changes in temperature. The invention includes the utilization of boron dopants in the core and cladding of optical waveguide devices such as Mach-Zehnder coupler devices and long period fiber gratings.

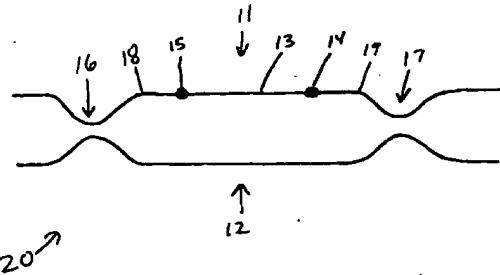


FIG. 1

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Description**FIELD OF THE INVENTION**

This invention relates to an optical waveguide device in which thermal spectral shifts are inhibited. In particular, this invention relates to athermalized optical waveguiding devices in which the light transmitting properties are insensitive to temperature variations and fluctuations.

BACKGROUND OF THE INVENTION

Optical waveguide devices, such as Mach-Zehnder coupler devices and long period fiber gratings, which are utilized with optical signals as transmitting filters, sensors, and wavelength multiplexing and demultiplexing devices provide good performance at standard room temperatures but exhibit thermal spectral shifts and related poor performance when used in environments where they are exposed to thermal variation and fluctuations in temperatures.

Mach-Zehnder coupler devices and long period fiber gratings can be used in narrow-band, multi-wavelength system applications as transmitting filters. In addition, Mach-Zehnder coupler devices can be used as sensors, and wavelength multiplexing and demultiplexing devices.

In a long period fiber grating a long period grating is formed in the fiber which functions to couple light between the fundamental mode propagating in the waveguide core and a guided cladding mode. Such gratings have an index modulation along the waveguiding axis of the fiber, and may be formed by writing with UV irradiation, etching, or other means of making periodic perturbations.

A Mach-Zehnder coupler device is a form of a Mach-Zehnder interferometer. In a Mach-Zehnder coupler device, a coupler splits light into two or more waveguides, typically optical fibers, which have different optical path lengths. A second coupler recombines this light after the light has traveled the different optical path lengths. In addition, Mach-Zehnder coupler devices may be used as demultiplexers and in sensor applications. Variations in the Mach-Zehnder coupler device family includes MultiClad™ Mach-Zehnder coupler devices, fused fiber Mach-Zehnder coupler devices and Mach-Zehnder lattice filters.

It has been found that the use of optical waveguide devices such as long period fiber gratings and Mach-Zehnder coupler devices is limited by their temperature dependence. In such devices, thermal spectral shifts of greater than 0.04 nm/°C at a transmitting wavelength of 1550 nm limit their application and usefulness in environments of differing temperature.

SUMMARY OF THE INVENTION

Accordingly, the present invention is directed to an optical waveguide that substantially obviates one or more of the problems due to limitations and disadvantages of the related art.

Additional features and advantages of the invention will be set forth in the description which follows, and in part will be apparent from the description, or may be learned by practice of the invention. The objectives and other advantages of the invention will be realized and attained by the compositions, structures, and methods particularly pointed out in the written description and claims hereof as well as the appended drawings.

To achieve these and other advantages and in accordance with the purpose of the invention, as embodied and broadly described, the invention provides an athermal optical waveguide device in which at least one light transmitting GeO_2 doped silica core is codoped with B_2O_3 to inhibit thermal spectral shifts induced by changes in temperature.

In another aspect, the invention includes an athermal optical waveguide doped with B_2O_3 .

A further aspect of the invention is to provide a method of athermalizing an optical waveguide by codoping the waveguide with B_2O_3 to neutralize the optical waveguide's thermal sensitivity.

It is to be understood that both the foregoing general description, and the following detailed description are exemplary and explanatory and are intended to provide further explanation of the invention as claimed.

The accompanying drawings are included to provide a further understanding of the invention and are incorporated in and constitute a part of this specification, illustrate several embodiments of the invention and, together with the description, serve to explain the principles of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 comprises a schematic representation of a Mach-Zehnder coupler device.

FIG. 2 is a plot of Temperature (°C) versus Peak Wavelength (nm) for a comparison model of a Mach-Zehnder coupler device.

FIG. 3 is a plot of Temperature (°C) versus Peak Wavelength (nm) for a Mach-Zehnder coupler device in accordance with the teachings of the present invention.

FIG. 4 is a plot of Temperature (°C) versus Peak Wavelength (nm) for a Mach-Zehnder coupler device in accordance with the teachings of the present invention.

FIG. 5 is a plot of Temperature (°C) versus Peak Wavelength (nm) for a Mach-Zehnder coupler device in accordance with the teachings of the present invention.

FIG. 6 is a plot of B_2O_3 Content (wt.%) versus Thermal Spectral Shift Coefficients (nm/°C) in accordance with the teachings of the present invention.

FIG. 7 is a plot of Temperature (°C) versus Peak

Wavelength (nm) for a long period fiber grating in accordance with the teachings of the present invention.

FIG. 8 is a plot of Temperature (°C) versus Peak Wavelength (nm) for a long period fiber grating in accordance with the teachings of the present invention.

FIG. 9 is a plot of Temperature (°C) versus Peak Wavelength (nm) for a long period fiber grating in accordance with the teachings of the present invention.

FIG. 10 is a plot of the B₂O₃ wt.% difference between the core and cladding of the long period fiber gratings versus the slope (nm/°C) of the long period fiber gratings at 20°C in accordance with the teachings of the present invention.

PRIOR ART

Prior art of possible relevance is provided separately.

DETAILED DESCRIPTION OF THE INVENTION

The optical waveguide device of the invention includes a germania doped silica core and a sufficient amount of B₂O₃ dopant to inhibit thermal spectral shifts.

Reference will now be made in detail to the present preferred embodiments of the invention, examples of which are illustrated in the accompanying drawings.

The athermal optical waveguide devices of the invention include optical waveguides, optical fibers, Mach-Zehnder coupler devices and long period fiber gratings which provide for the transmission of light through a silica core. The athermal optical waveguide devices of the invention are useable over a broad temperature range as transmitting filters, demultiplexers, multiplexers, and sensors.

The optical waveguide devices of the invention include an optical waveguiding silica core doped with an index raising dopant and a sufficient amount of B₂O₃ codopant to inhibit thermal spectral shifts in the device when the device is subjected to changes in temperature. The preferred index raising dopant of the invention is GeO₂.

The optical waveguide devices of the invention include transmitting filters. The transmitting filters of the invention include long period fiber gratings and Mach-Zehnder interferometers. A Mach-Zehnder coupler device of the invention utilizes a Mach-Zehnder interferometer in which a mode of light guided by a first waveguiding core is coupled to a second waveguiding core. The long period fiber grating transmitting filters of the invention provide for a mode of light propagating in a waveguiding core to couple with a mode in the cladding of the waveguide.

The optical waveguide devices of the invention include a silica core which has a B₂O₃:GeO₂ ratio ranging from 1 wt.% B₂O₃ : 3 wt.% GeO₂ to 1 wt.% B₂O₃ : 10 wt.% GeO₂, more preferably ranging from 1 wt.% B₂O₃ : 4 wt.% GeO₂ to 1 wt.% B₂O₃ : 6 wt.% GeO₂. A

preferred silica core has a 1 wt.% B₂O₃ : 5 wt.% GeO₂ ratio.

The athermalized optical waveguide includes a silica glass composition which has a dopant content of B₂O₃ and one or more index raising dopants, wherein the dopant content has a B₂O₃ : index raising dopant molar ratio ranging from 1 mole B₂O₃ : 1.5 moles of index raising dopants to 1 mole B₂O₃ : 8 moles of index raising dopants. A preferred dopant content range is from 1 mole B₂O₃ : 2 moles of index raising dopants to 1 mole B₂O₃ : 7 moles of index raising dopants. A more preferred dopant content range is from 1 mole B₂O₃ : 3 moles of index raising dopants to 1 mole B₂O₃ : 4 moles of index raising dopants.

The method of making the optical waveguide devices of the invention includes the method of athermalizing the optical waveguide device to inhibit thermal spectral shift. This method includes forming a silica waveguide core and doping the core with an index raising dopant, preferably GeO₂. This method further includes the step of codoping the core with B₂O₃. Codoping with B₂O₃ provides for a sufficient amount of B₂O₃ wherein the waveguide device exhibits a thermal spectral shift of less than 0.04 nm/°C at 1550 nm when subjected to a temperature variation. The core is codoped with a sufficient amount of B₂O₃ to neutralize the core's thermal sensitivity.

This method may further include the step of forming a Mach-Zehnder interferometer with the codoped silica waveguide core.

This method can also include the step of cladding the codoped waveguide core to form an optical waveguide. Such a cladding can be comprised of silica or B₂O₃ doped silica. An additional step of the invention is to write a long period grating in the optical waveguide.

The invention includes the method of reducing the temperature dependence of optical waveguide devices. This method includes the steps of forming a waveguide core, doping the waveguide core with an index raising dopant having a positive thermal spectral shift slope, and codoping the waveguide core with a second dopant having a negative thermal spectral shift slope. Codoping the core includes the step of neutralizing the positive thermal spectral shift slope of the core with the negative thermal spectral shift slope of the second dopant. Said second dopant is preferably boron, which is an index depressing dopant. The step of codoping can further comprise codoping with an amount of the second dopant so that the molar ratio of the second dopant to the index raising dopant is in the range of 1 mole of the second dopant to 2-4 moles of the index raising dopant.

This method further includes the steps of cladding the waveguide core and fabricating an optical waveguide device from the cladded waveguide core. Such a cladding may include the use of a glass doped with the second dopant.

The invention includes the method of making an optical waveguide with a reduced temperature depend-

ence by doping a waveguide with an index altering dopant which allows light to be guided and doping the waveguide with a temperature dependence liberating dopant which reduces the temperature dependence of the waveguide.

The athermalized optical waveguide of the invention includes a core and a cladding wherein the core and the cladding have a difference in B_2O_3 dopant content ranging from 1 to 5 wt.% B_2O_3 , with a preferred difference ranging from 2 to 4 wt.% B_2O_3 .

U.S. Patent Number 5,295,205 to Miller et al., the disclosure of which is hereby incorporated by reference, discloses a Mach-Zehnder coupler device and a method of making such devices.

One embodiment of the invention comprises a MultiClad™ Mach-Zehnder coupler device which utilizes a first optical fiber having a silica core doped with GeO_2 which is codoped with B_2O_3 in order to inhibit thermal spectral shifts in the Mach-Zehnder coupler device caused by a change in temperature. The Mach-Zehnder coupler device of the invention further includes a second optical fiber having a B_2O_3 dopant concentration different from the first optical fiber. Such an embodiment provides a Mach-Zehnder coupler device that has a reduced sensitivity to temperature variations.

A number of MultiClad™ Mach-Zehnder coupler devices of the type having two arms of equal physical path lengths but differing fundamental mode effective indices, as illustrated in FIG. 1 by Mach-Zehnder coupler device 20, were made and analyzed at temperatures ranging from -40°C to 125°C. In FIG. 1, spliced Mach-Zehnder coupler device arm 11 has the same physical path length as unspliced Mach-Zehnder coupler device arm 12. First coupler 16 splits light into spliced coupler arm 11 and unspliced coupler arm 12. Spliced coupler arm 11 has an optical path length different from the optical path length of unspliced coupler arm 12 due to the insertion of optical path changing fiber segment 13. Second coupler 17 recombines the split light propagating through coupler arms 11 and 12. In Mach-Zehnder coupler device 20 light is selectively transmitted by the Mach-Zehnder interferometer by being split into two differing optical path lengths and then recombined. The invention may also be utilized with other forms of Mach-Zehnder interferometers and coupler devices, for example Mach-Zehnder coupler devices with more than two arms, Mach-Zehnder coupler devices with unequal coupler splitting, and Mach-Zehnder coupler devices having arms of different physical path lengths.

A comparison model of a Mach-Zehnder coupler device was made as shown in FIG. 1 with commercially available optical fibers. Experimental Mach-Zehnder coupler devices of the invention were made as shown in FIG. 1 and compared with the comparison model. In the comparison model the Mach-Zehnder coupler device's unspliced coupler arm 12 was made from commercially available Corning SMF-28™ single mode optical fiber.

Spliced coupler arm 11, which has the same physical path length as unspliced coupler arm 12, was made by splicing in a segment of commercially available Corning CS-980™ single mode optical fiber. Corning SMF-28™ single mode optical fiber has a core-clad $\Delta = 0.35\%$ with a cutoff of 1260 nm and a GeO_2 doped silica core, which is doped with about 6-10 wt.% GeO_2 . Corning CS-980™ single mode optical fiber has a core-clad $\Delta = 1.0\%$ with a cutoff of 950 nm and a GeO_2 doped silica core, which is doped with about 18-20 wt.% GeO_2 . In this comparison model spliced in fiber segment 13, a 10 mm segment of Corning CS-980™ single mode optical fiber was spliced between Corning SMF-28™ single mode optical fiber segments 18 and 19 at splicings 15 and 14. The thermal spectral shift of this Mach-Zehnder comparison model coupler was analyzed using an optical spectrum analyzer and an erbium ASE light source. The performance of this comparison model at a variety of temperatures in the range of -40°C to 125°C was tested using a thermal chamber. The thermal spectral shift of the comparison model is shown in FIG. 2 which is a plot of Peak Wavelength (nm) versus Temperature (°C). From FIG. 2 it was determined that this comparison model had a thermal spectral shift coefficient of 0.043 nm/°C. Virtually identical thermal spectral shift coefficients were measured for other comparison models which had a variety of GeO_2 core dopant levels, arm lengths and fiber types such as Corning SMF-28™ and CS-980™ optical fibers with different cutoffs, optical fibers with a TiO_2 doped silica core, and a step index optical fiber having a $\Delta = 2.0\%$ with a cutoff of 1210 nm.

Experimental Mach-Zehnder coupler device No. 1 was made with the configuration of Mach-Zehnder coupler device 20 of FIG. 1 and in the same set up and manner as the comparison model except that spliced in fiber length 13 was not Corning CS-980™ optical fiber. The spliced in fiber length 13 of experimental coupler No. 1 was a 10 mm section of an experimental fiber having an optical waveguiding silica core doped with 25 wt.% GeO_2 and codoped with 14 wt.% B_2O_3 . (16.5 mole % GeO_2 , 13.7 mole % B_2O_3 , and 69.8 mole % SiO_2) The thermal spectral shift of experimental coupler No. 1 was analyzed in the same manner as the comparison model. The thermal spectral shift of experimental coupler No. 1 is shown in FIG. 3 which is a plot of Peak Wavelength (nm) versus Temperature (°C). From FIG. 3 it was determined that experimental coupler No. 1 had a thermal spectral shift coefficient of -0.204 nm/°C.

Experimental Mach-Zehnder coupler device No. 2 was made with the configuration of Mach-Zehnder coupler device 20 of FIG. 1 and in the same manner and setup as the comparison model except that spliced in fiber length 13 was a fiber segment comprised of a 7.757 mm length of Corning CS-980™ single mode optical fiber, as used in the comparison model, spliced with a 1.158 mm length of experimental fiber having a silica core doped with 25 wt.% GeO_2 and codoped with 14 wt.% B_2O_3 , as used in experimental coupler No. 1. This

combination of spliced fiber for experiment coupler No. 2 was chosen so as to achieve a net zero thermal spectral shift coefficient. The thermal spectral shift of experimental coupler No. 2 was analyzed in the same manner as the comparison model. The thermal spectral shift of experimental coupler No. 2 is shown in FIG. 4 which is a plot of Peak Wavelength (nm) versus Temperature (°C). From FIG. 4 it was determined that experimental coupler No. 2 had a thermal spectral shift coefficient of -0.008 nm/°C.

Experimental Mach-Zehnder coupler device No. 3 was made with the configuration of Mach-Zehnder coupler device 20 of FIG. 1 and in the same manner and set up as the comparison model except that spliced in fiber length 13 was a 10 mm section of an experimental fiber having an optical waveguiding silica core doped with 15 wt.% GeO₂ and codoped with 5 wt.% B₂O₃. (9.3 mole % GeO₂, 4.6 mole % B₂O₃, 86.1 mole % SiO₂) This experimental codoped fiber was fabricated by the outside vapor deposition process using a flame hydrolysis apparatus with appropriate silica, GeO₂ dopant, and B₂O₃ codopant feedstock levels to result in this core which was cladded with silica. The thermal spectral shift of experimental coupler No. 3 was analyzed in the same manner as the comparison model. The thermal spectral shift of experimental coupler No. 3 is shown in FIG. 5 which is a plot of Peak Wavelength (nm) versus Temperature (°C). From FIG. 5 it was determined that experimental coupler No. 3 had a thermal spectral shift coefficient of -0.07 nm/°C.

Hypothetical experimental Mach-Zehnder coupler device No. 4 would be fabricated from a unitary spliced in fiber length 13 to avoid the complications of a spliced in fiber length 13 made from a GeO₂ doped fiber and a GeO₂ doped B₂O₃ codoped fiber as in experimental Mach-Zehnder coupler device No. 2 to achieve an athermalized optical waveguide device with a nearly zero thermal spectral shift coefficient. FIG. 6 shows the thermal spectral shift coefficients of the comparison model (.043 nm/°C at 0 wt.% B₂O₃), experimental coupler No. 3 (-0.07 nm/°C at 5 wt.% B₂O₃), and experimental coupler No. 1 (-0.215 nm/°C at 14 wt.% B₂O₃). FIG. 6 is a plot of the thermal spectral shift coefficients of these couplers in nm/°C versus the B₂O₃ content of these couplers in wt.%. FIG. 6 provides an optimal level of boron codoping of GeO₂ doped silica cores to achieve temperature independence. Linear interpolation of the data as shown in FIG. 6 gives an optimal level of boron codoping of 15 wt.% GeO₂ doped silica cores of approximately 1.7± 0.6 wt.% B₂O₃ (9.3 mole % GeO₂, 1.6 mole % B₂O₃, 89.1 mole % SiO₂) to achieve a thermal spectral shift coefficient of zero nm/°C. This optimal and sufficient B₂O₃ codoping concentration level is useable with a broad range of GeO₂ core doping levels since the thermal spectral shift coefficients of silica core waveguides doped only with germania are independent of the germania doping levels within a broad range of about 10 wt.% to 40 wt.% GeO₂.

Hypothetical experimental coupler No. 4 would be fabricated with the same configuration as FIG. 1 and in the same manner and set up of the comparison model except that spliced in fiber length 13 would be a 10 mm section of an experimental fiber having an optical waveguiding silica core doped with 1 wt.% to 40 wt.% of GeO₂, preferably 15 wt.% to 25 wt.% of GeO₂ and codoped with 1 wt.% to 4 wt.% B₂O₃, preferably 1.1 wt.% to 2.3 wt.% of B₂O₃. This experimental codoped fiber could be fabricated by the outside vapor deposition process using a flame hydrolysis apparatus with appropriate silica, GeO₂ dopant, and B₂O₃ codopant feedstock levels, and also by other processes, to result in such a core, which could be cladded with silica.

An additional embodiment of the invention includes the fabrication of a long period fiber gratings from an optical fiber having a silica core codoped with B₂O₃. Long period fiber gratings of the invention were formed using UV Excimer lasers with appropriate energies, exposure times, and grating periods.

A comparison model of a long period fiber grating was fabricated from commercially available Corning SMF-DST™, which is a dispersion shifted fiber having a dispersion shifting core profile with a peak of about 18-20 wt.% GeO₂. Long period gratings were written in this fiber using a laser with 550mJ pulses at 248 nm and at an 8 Hz repetition rate for 6 minutes. Prior to writing this long period grating, the fiber was placed in a hydrogen atmosphere at 1550 psi for one week to increase the photosensitivity of the fiber. This resulted in the comparison model having a strong grating with only 9% power transmission. The thermal spectral shift of this comparison model long period fiber grating was analyzed in a similar manner as the Mach-Zehnder coupler devices. This comparison model long period fiber grating had a thermal spectral shift coefficient of 0.044 nm/°C.

Experimental long period fiber grating No. 1 was fabricated from an experimental optical fiber having a silica core doped with 15 wt.% GeO₂ and codoped with 5 wt.% B₂O₃. This experimental codoped fiber core was cladded with a silica cladding having approximately 0.8 wt.% B₂O₃. Experimental long period fiber grating No. 1 was hydrogen treated then written in the same manner as the comparison model. This resulted in a weaker grating than the comparison model. Experimental long period fiber grating No. 1 had a 62% power transmission indicating that this experimental optical fiber was less photosensitive than the conventional dispersion-shifted fiber used in the comparison model. The thermal spectral shift of this long period fiber grating was analyzed in a similar manner as the Mach-Zehnder coupler devices and the long period fiber grating comparison model. The thermal spectral shift of experimental long period fiber grating No. 1 is shown in FIG. 7 which is a plot of Peak Wavelength (nm) versus Temperature (°C). From FIG. 7 it was determined that experimental long period fiber grating No. 1 had a thermal spectral shift coefficient of -0.028 nm/°C. It is believed that experimental

long period fiber grating No. 1 is more athermalized than experimental Mach-Zehnder coupler device No. 3 which used the same fiber core compositions of GeO₂ and B₂O₃ due to the presence of boron in the cladding.

Experimental long period fiber grating No. 2 was fabricated from an experimental optical fiber having a silica core doped with 15 wt.% GeO₂ and codoped with 2 wt.% B₂O₃. This experimental codoped fiber core was cladded with a silica cladding having 1 wt.% B₂O₃. Experimental long period fiber grating No. 2 was hydrogen treated then written in the same manner as the comparison model. The thermal spectral shift of this long period fiber grating was analyzed in a similar manner as the Mach-Zehnder coupler devices and the long period fiber gratings. The thermal spectral shift of experimental long period fiber grating No. 2 is shown in FIG. 8 which is a plot of Peak Wavelength (nm) versus Temperature (°C). From FIG. 8 it was determined that experimental long period grating No. 2 had a thermal spectral shift coefficient of 0.026 nm/°C.

Experimental long period fiber grating No. 3 was fabricated from an experimental optical fiber having a silica core doped with 17 wt.% GeO₂ and codoped with 3.3 wt.% B₂O₃. This experimental codoped fiber core was cladded with a silica cladding having 0.9 wt.% B₂O₃. Experimental long period fiber grating No. 3 was hydrogen treated then written in the same manner as the comparison model. The thermal spectral shift of this long period fiber grating was analyzed in a similar manner as the Mach-Zehnder coupler devices and the long period fiber gratings. The thermal spectral shift of experimental long period fiber grating No. 3 is shown in FIG. 9 which is a plot of Peak Wavelength (nm) versus Temperature (°C). From FIG. 9 it was determined that experimental long period grating No. 3 had a thermal spectral shift coefficient of 0.004 nm/°C.

FIG. 10 is a plot of the thermal spectral shift coefficients of the experimental long period fiber gratings and the comparison model versus the B₂O₃ wt.% difference between the core and the cladding for each of these long period fiber gratings. As shown in FIG. 10, the comparison model had a thermal spectral shift coefficient of 0.044 nm/°C and 0 B₂O₃ wt.% difference between its core and cladding in that it was not doped with boron. Experimental long period fiber grating No. 1 had a thermal spectral shift coefficient of -0.028 nm/°C and a 4.2 B₂O₃ wt.% difference between its core of 5 wt.% B₂O₃ and cladding of 0.8 wt.% B₂O₃. Experimental long period fiber grating No. 2 had a thermal spectral shift coefficient of 0.026 nm/°C and a 1 B₂O₃ wt.% difference between its core of 2 wt.% B₂O₃ and cladding of 1 wt.% B₂O₃. Experimental long period fiber grating No. 3 had a thermal spectral shift coefficient of 0.004 nm/°C and a 2.4 B₂O₃ wt.% difference between its core of 3.3 wt.% B₂O₃ and cladding of 0.9 wt.% B₂O₃. As shown in FIG. 10 it is preferred to have an optical fiber with a difference in B₂O₃ wt.% between the core and the cladding of about 2 to 3.2 to achieve a thermal spectral shift

coefficient from 0.01 to -0.01 nm/°C. An optimized athermalized long period fiber grating would have a difference of about 2.6 wt.% between the B₂O₃ concentrations of the core and the cladding.

Hypothetical experimental long period fiber grating No. 4 would be fabricated from an experimental optical fiber having a difference of 2.6 between the B₂O₃ wt % of the core and the cladding, such as the silica core of experimental long period fiber grating No. 3 being clad- ded with a silica cladding having 0.7 wt.% B₂O₃.

An optical waveguiding silica core having about 1 wt.% to 40 wt.% of GeO₂, preferably 15 wt.% to 25 wt.% of GeO₂ and codoped with 1 wt.% to 5 wt.% of B₂O₃, preferably 2 wt.% to 4.5 wt.% of B₂O₃, and cladded with a silica cladding having from about 0 wt.% to 2 wt.% of B₂O₃, preferably 0.6 to 1 wt.% of B₂O₃ is preferred to achieve an athermalized temperature independent optical waveguide device, such as long period fiber grating. In addition, to achieve such an athermalized tempera- ture independent optical waveguide device, particularly a long period fiber grating it is important that the differ- ence in B₂O₃ concentration between the core and the cladding be in the range of about 2 to 3.2 wt.%, prefera- bly 2.2 to 3 wt.%, as disclosed in FIG. 10.

As with the long period fiber gratings, B₂O₃ could be used in the silica cladding surrounding the core of an optical waveguide fiber in a Mach-Zehnder coupler device. With the optical waveguide fibers of the invention, the actual B₂O₃ concentration in the cladding has a relatively small effect compared to the difference of B₂O₃ concentration in the core and the cladding. In addition, Mach-Zehnder coupler devices could be fabri- cated that have more than one of the fibers doped with boron.

The experimental fibers of the invention were fabri- cated with a flame hydrolysis apparatus using the out- side vapor deposition method to form these optical waveguide silica cores doped with GeO₂ and codoped with B₂O₃. Appropriate silica, GeO₂ dopant, and B₂O₃ codopant vaporous feedstock levels are delivered to the flame hydrolysis burner to provide for the particular dopant and codopant concentration levels. Similarly, appropriate silica and B₂O₃ vaporous feedstock levels are delivered to the flame hydrolysis burner to provide for the particular B₂O₃ concentrations in the silica clad- ding glass used in the long period fiber gratings.

It is believed that this inventive codoping of a GeO₂ doped silica core with B₂O₃ beneficially changes the thermal variability of the index of refraction of the waveguiding core, along with the potentially advan- tageous altering of the waveguide dispersion effects and stress-optic effects of the optical fiber.

The waveguide core and optical fiber composition of the invention is particularly sensitive and dependent on the B₂O₃ dopant concentration. In the inventive waveguide core composition, GeO₂ is utilized as a dopant to advantageously alter the index of refraction of the silica core in order to achieve a beneficial Δ for guid-

ing the light and to provide photosensitivity in the long period fiber gratings. The B_2O_3 is utilized as a codopant to advantageously alter the thermal variability of refractive index of the silica core. A B_2O_3 to GeO_2 concentration ratio in the range of 1 wt.% B_2O_3 : 3 wt.% GeO_2 to 1 wt.% B_2O_3 : 10 wt.% GeO_2 will provide a beneficial thermal variability which inhibits and reduces thermal spectral shifts while providing an appropriate waveguide core refractive index. The preferred B_2O_3 to GeO_2 concentration ratio is in the range of 1 wt.% B_2O_3 : 4 wt.% GeO_2 to 1 wt.% B_2O_3 : 6 wt.% GeO_2 . The most preferred B_2O_3 to GeO_2 concentration ratio of the invention is 1 wt.% B_2O_3 : 5 wt.% GeO_2 .

In view of obtaining an advantageous core to cladding B_2O_3 concentration difference, the preferred core composition is from 1-40 wt.% GeO_2 and 0.5-15 wt.% B_2O_3 . The more preferred core composition is from 4-25 wt.% GeO_2 and 0.75-7 wt.% B_2O_3 . The most preferred core composition is from 6-20 wt.% GeO_2 and 1-3 wt.% B_2O_3 .

Also in view that in the invention GeO_2 is used as an index raising altering dopant to guide light waves and that B_2O_3 is used as a negative thermal spectral shift dopant to neutralize the positive thermal spectral shift of the index raising dopant and silica composition, the preferred molar ratio is 1 mole of B_2O_3 to 2-4 total moles of index raising dopants which includes dopants such as germanium and phosphorous. The most preferred molar ratio of negative thermal spectral shift dopants to the total moles of index raising dopants is 1 mole of B_2O_3 to 3 total moles of index raising dopants to achieve an athermalized optical waveguide.

It will be apparent to those skilled in the art that various modifications and variations can be made in the composition and method of the present invention without departing from the spirit or scope of the invention. Thus, it is intended that the present invention cover the modifications and variations of the invention provided they come within the scope of the appended claims and their equivalents.

Claims

1. An optical waveguide device comprised of an optical waveguiding silica core doped with an index raising dopant, wherein said waveguiding core is codoped with a sufficient amount of B_2O_3 to inhibit thermal spectral shifts in the device caused by a change in the temperature of said device
2. An optical waveguide device as claimed in claim 1 wherein said index raising dopant is GeO_2 .
3. An optical waveguide device as claimed in claim 1 wherein said waveguide device comprises a transmitting filter.
4. An optical waveguide device as claimed in claim 1,
5. An optical waveguide device as claimed in claim 1, further comprising a cladding, wherein a mode propagating in said core is coupled to a mode in said cladding.
- 10 6. An athermalized optical waveguide device comprised of a silica glass with a dopant content consisting essentially of GeO_2 and B_2O_3 .
- 15 7. An optical waveguide device as claimed in claim 2 or 6 wherein said silica core has a B_2O_3 : GeO_2 ratio ranging from
1 wt.% B_2O_3 : 3 wt.% GeO_2 to
1 wt.% B_2O_3 : 10 wt.% GeO_2 , or
- 20 8. a B_2O_3 : GeO_2 ratio ranging from
1 wt.% B_2O_3 : 4 wt.% GeO_2 to
1 wt.% B_2O_3 : 6 wt.% GeO_2 , or
- 25 9. wherein said silica core has a B_2O_3 : GeO_2 ratio of 1 wt.% B_2O_3 : 5 wt.% GeO_2 , or a B_2O_3 : GeO_2 molar ratio ranging from 1 mole B_2O_3 : 1.5 moles GeO_2 to 1 mole B_2O_3 : 8 moles GeO_2 .
- 30 10. An optical waveguide device as claimed in claim 4 or 5 wherein said device is comprised of a Mach-Zehnder interferometer, or of a long period fiber grating.
- 35 11. An athermalized optical waveguide comprised of a silica glass with a dopant content comprised of B_2O_3 and a plurality of index raising dopant molecules.
- 40 12. A waveguide as claimed in claim 9 wherein said dopant content has a B_2O_3 : index raising dopant molar ratio ranging from 1 mole B_2O_3 : 2 moles of index raising dopant molecules to 1 mole B_2O_3 : 7 moles of index raising dopant molecules, or has a B_2O_3 : index raising dopant molar ratio ranging from 1 mole B_2O_3 : 3 moles of index raising dopant molecules to 1 mole B_2O_3 : 4 moles of index raising dopant molecules.
- 45 13. A method of athermalizing an optical waveguide device to inhibit thermal spectral shifts, said method comprising:
(a) forming a silica waveguide core;
(b) doping said silica waveguide core with GeO_2 ; and
(c) codoping said silica waveguide core with B_2O_3 .
- 50 14. A method of athermalizing an optical waveguide device to inhibit thermal spectral shifts, said method comprising:
(a) forming a silica waveguide core;
(b) doping said silica waveguide core with GeO_2 ; and
(c) codoping said silica waveguide core with B_2O_3 .
- 55 15. A method of athermalizing an optical waveguide device to inhibit thermal spectral shifts, said method comprising:
(a) forming a silica waveguide core;
(b) doping said silica waveguide core with GeO_2 ; and
(c) codoping said silica waveguide core with B_2O_3 .

12. The method of athermalizing an optical waveguide device as claimed in claim 11 wherein the step of codoping the silica waveguide core with B₂O₃ comprises the step of codoping said silica waveguide core with a sufficient amount of B₂O₃ wherein said waveguide device exhibits a thermal spectral shift of less than 0.04 nm°C at 1550 nm. 5

13. The method of athermalizing an optical waveguide device as claimed in claim 11, wherein said method further comprises the steps of: 10

codoping said GeO₂ doped silica waveguide core with a sufficient amount of B₂O₃ to neutralize said GeO₂ doped silica waveguide core's thermal sensitivity. 15

14. The method of athermalizing an optical waveguide device as claimed in claim 11, further comprising the steps of: 20

forming a Mach-Zehnder interferometer with said codoped silica waveguide core, or cladding said codoped waveguide core to form an optical waveguide; and writing a long period grating in said formed optical waveguide. 25

15. The method of athermalizing an optical waveguide device as claimed in claim 11 wherein said method further comprises the step of cladding said codoped waveguide core with a B₂O₃ doped silica. 30

16. A method of reducing the temperature dependance of an optical waveguide device, said method comprising: 35

a) forming a waveguide core;
 b) doping the waveguide core with an index raising dopant having a positive thermal spectral shift slope; 40
 c) codoping said waveguide core with a second dopant having a negative thermal spectral shift slope;
 d) cladding said waveguide core; and 45
 e) fabricating an optical waveguide device from said cladded codoped waveguide core. 50

17. The method of claim 16, wherein said step of codoping further comprises the step of neutralizing the positive thermal spectral shift slope of the waveguide device with the second dopant, or of codoping said waveguide core with an index depressing dopant, or codoping said waveguide core with boron, or codoping with an amount of said second dopant so that the molar ratio of said second dopant to said index raising dopant is in the range of 1 mole of said second dopant to 2-4 moles 55

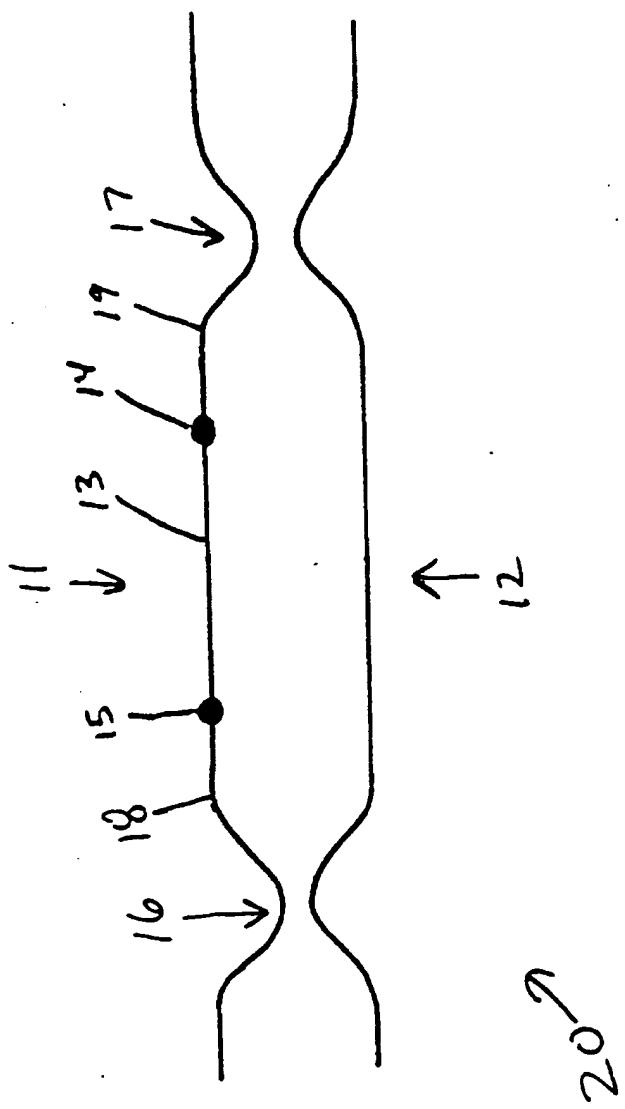


FIG 6. 1

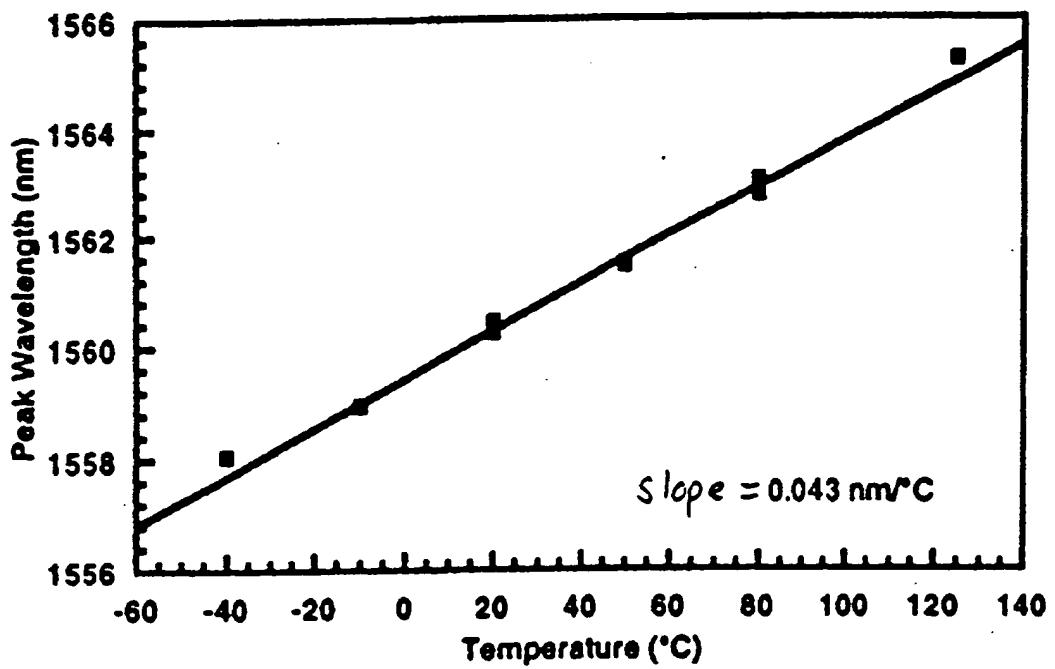


FIG. 2

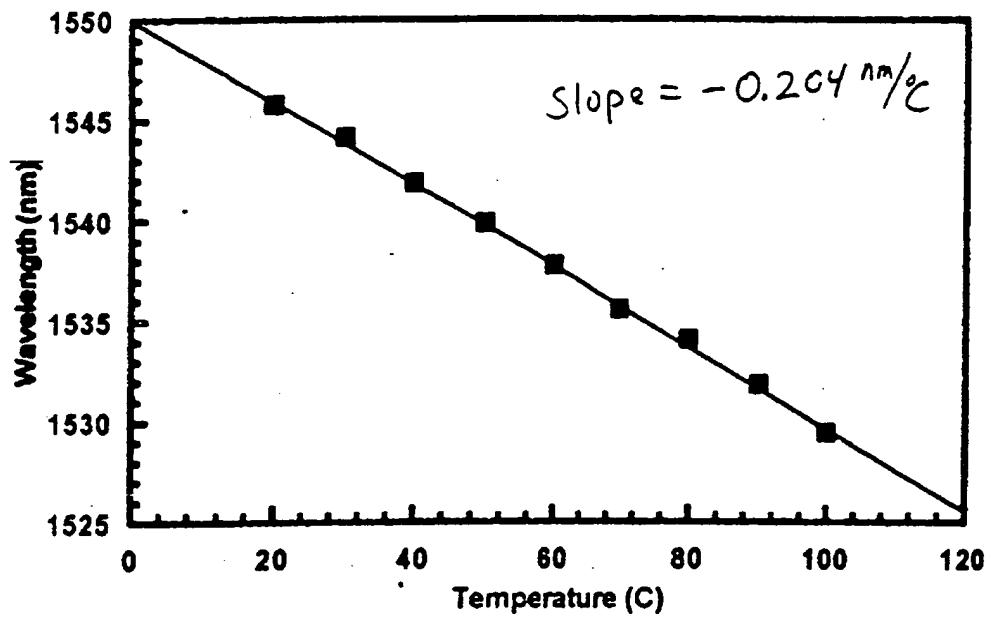


FIG. 3

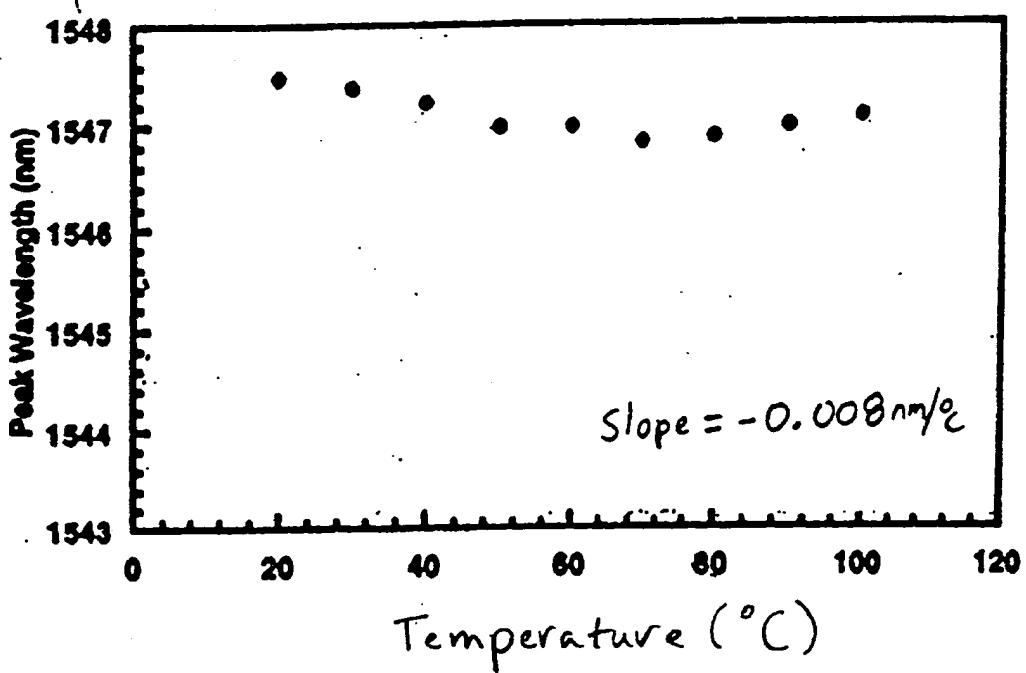


FIG. 4

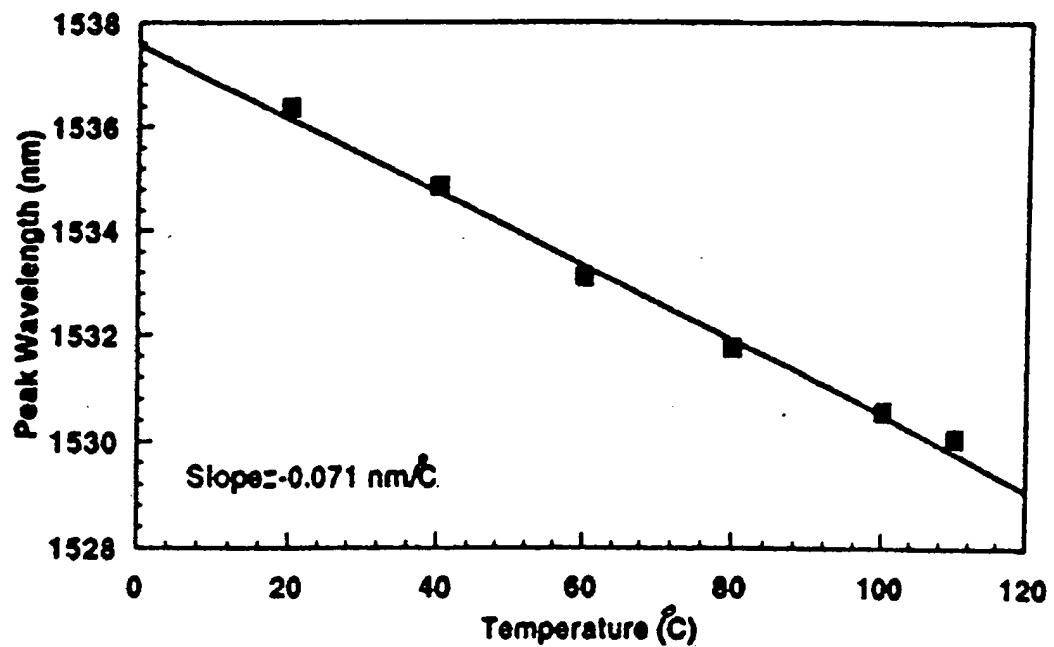


FIG. 5

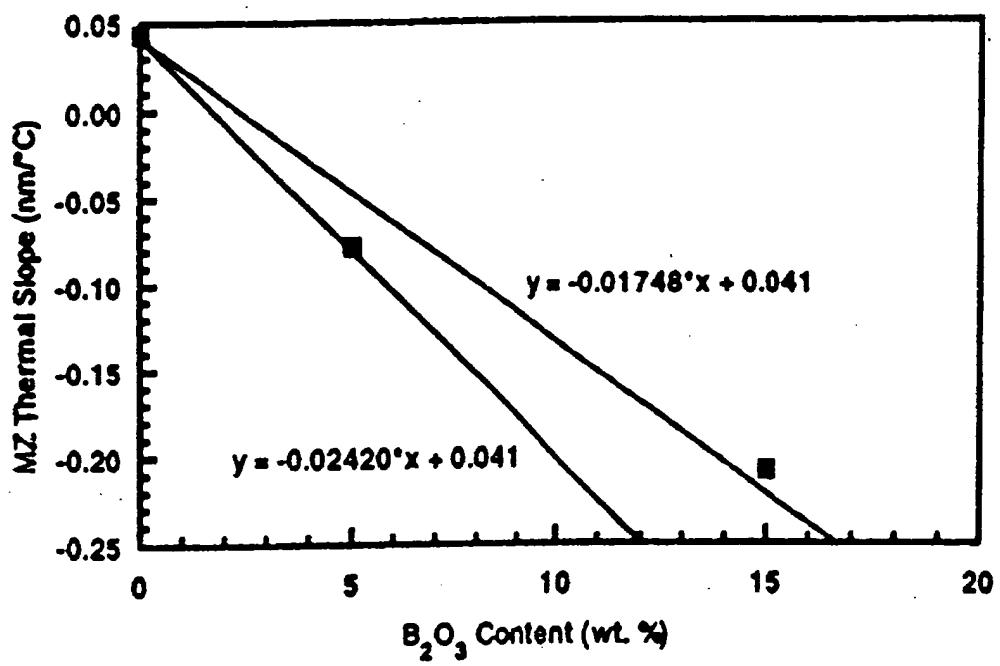


FIG. 6

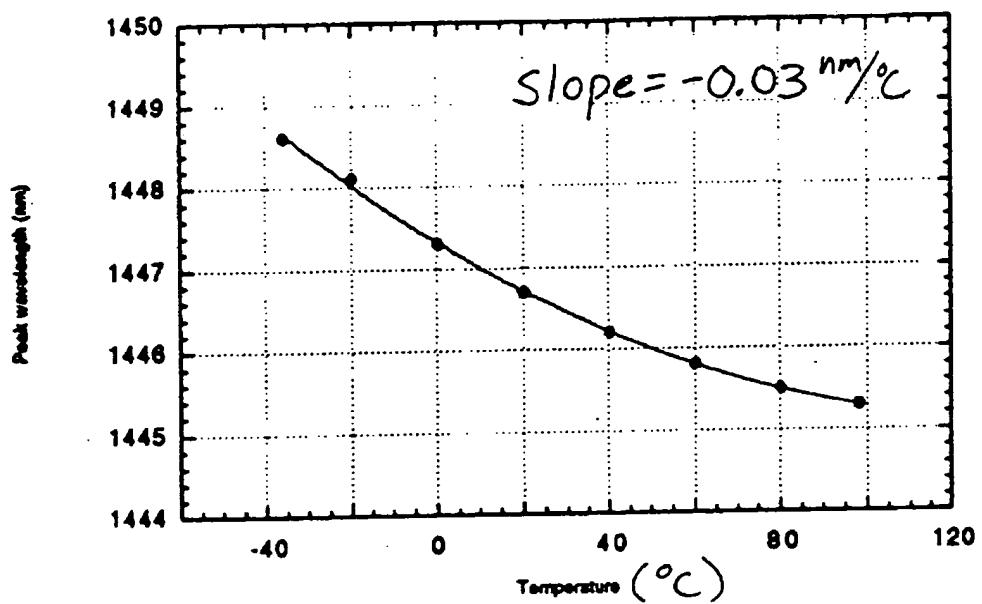
5 wt.% B₂O₃ in core

FIG. 7

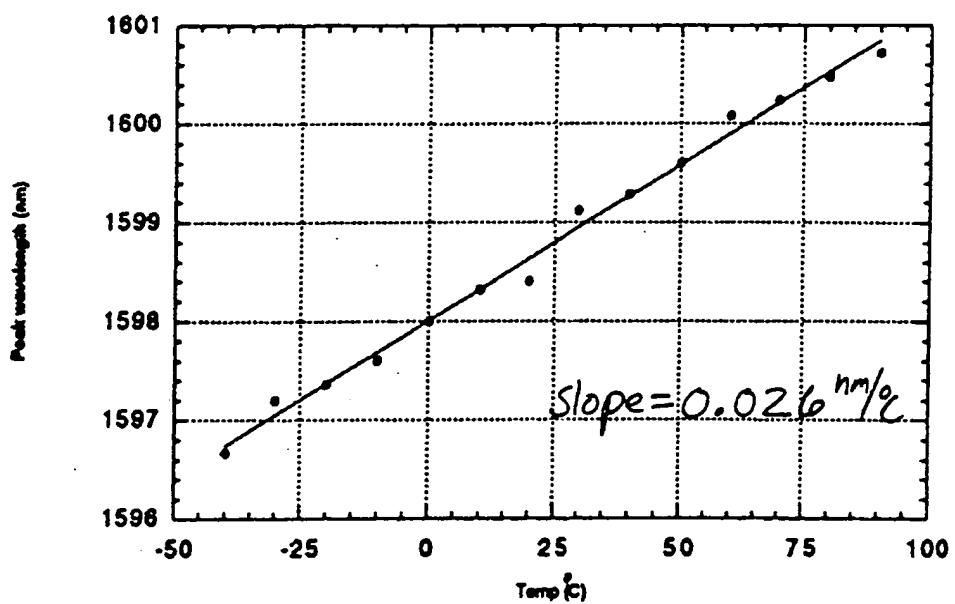
2 wt.% B₂O₃ in core

FIG. 8

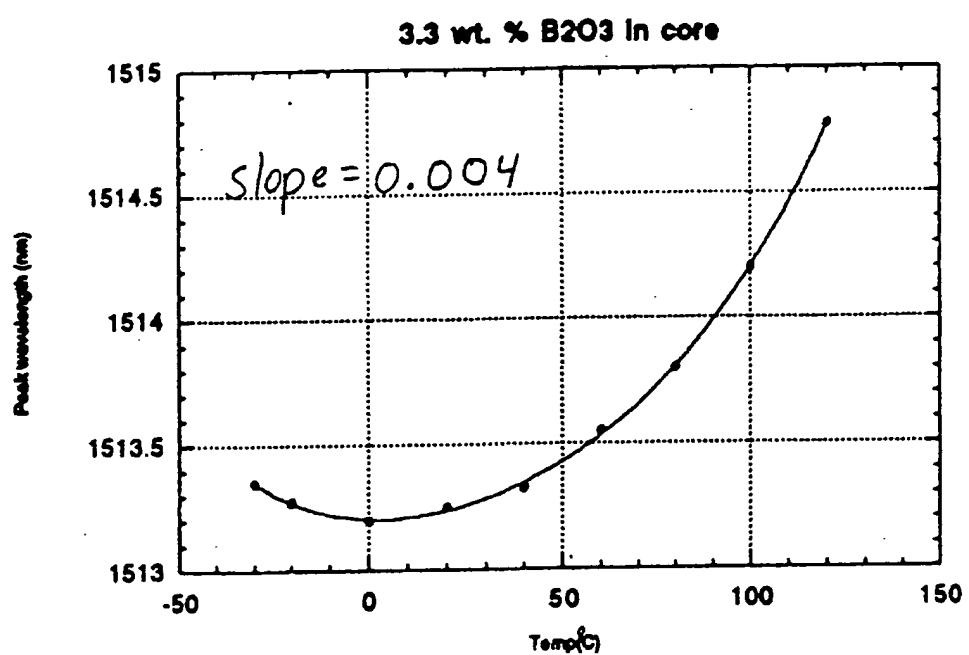


FIG. 9

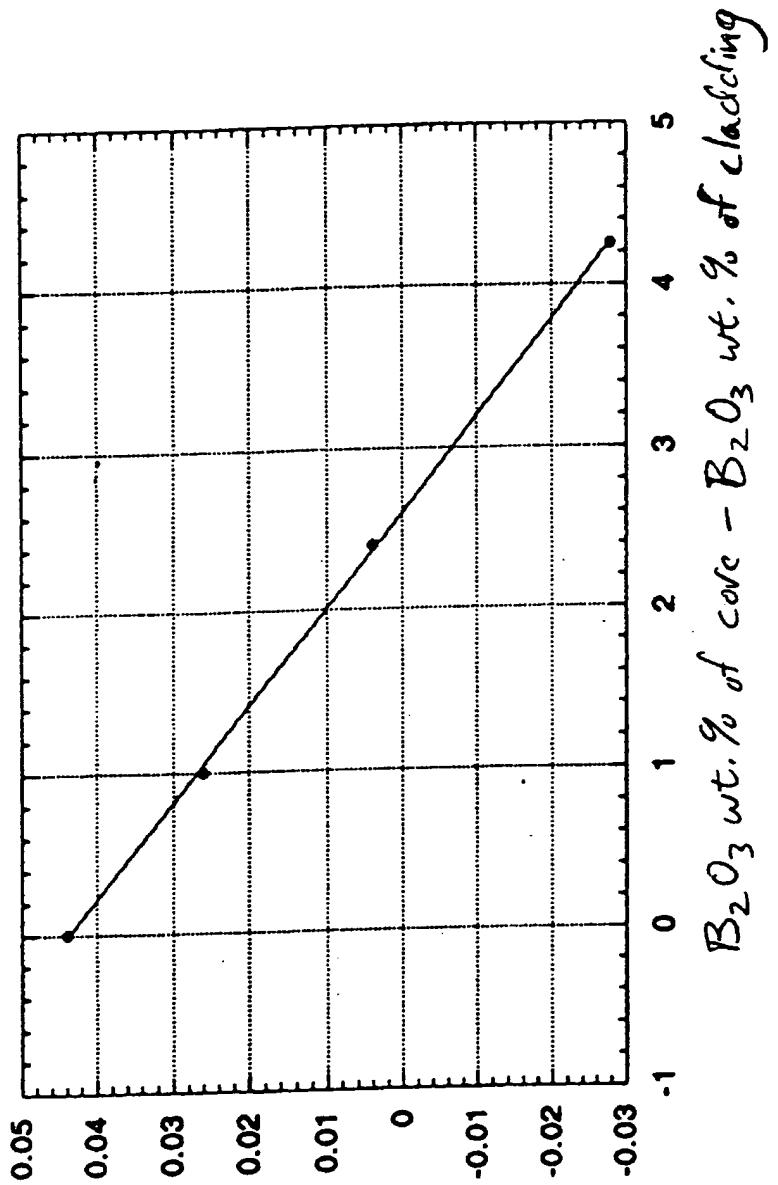


FIG. 10

long period fiber growth slope (nm/C)



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Application Number

EP 97 12 1532

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Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
P, X	EP 0 800 098 A (FUJIKURA LTD) 8 October 1997 * page 4, line 3 - page 5, line 27 * * page 13, line 20 - line 24 * * page 13, line 54 - line 55 * ---	1-20	C03B19/14 C03C13/04 G02B6/16 G02B6/26
X	WILLIAMS D L ET AL: "TEMPERATURE STABLE 1.3 MUM LASER WITH BRAGG FIBRE GRATING EXTERNAL CAVITY FOR ACCESS NETWORKS" PROCEEDINGS OF THE EUROPEAN CONFERENCE ON OPTICAL COMMUNICATION (EC, MONTREUX, SEPT. 12 - 16, 1993 REGULAR PAPERS, vol. VOL. 2, no. CONF. 19, 1993, SWISS ELECTROTECHNICAL ASSOCIATION, pages 209-212, XP000655040 * the whole document * ---	1-6,8, 11-14, 16,17,20	
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-/--			
The present search report has been drawn up for all claims			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	25 March 1998	Van Bommel, L	
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			



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EUROPEAN SEARCH REPORT

Application Number
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DOCUMENTS CONSIDERED TO BE RELEVANT		Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
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The present search report has been drawn up for all claims			
Place of search		Date of completion of the search	Examiner
THE HAGUE		25 March 1998	Van Bommel, L
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